GRAIN PROTEIN-BASED FORMULATIONS AND METHODS OF USING SAME

FIELD OF THE INVENTION

[0001] The present invention broadly concerns a grain protein formulation having an improved shelf life. More particularly, protein resin formulations may be manufactured in pelletized form well in advance of final production processing that converts the pellets into shaped articles of manufacture, for example, by injection molding, extrusion or other forming equipment. Shelf life of the protein resin formulations is enhanced by the addition of shelf stabilizing agents, such as hydrolyzed proteins, hydrolyzed protein derivatives, and hydrolyzed protein/hydrolyzed protein derivative-emulsifier complexes. The articles of manufacture from such resins may be pet chew treats, edible products, and biodegradable articles.

BACKGROUND OF THE INVENTION

[0002] Petroleum-based synthetic resins have achieved widespread use in the fabrication of a multitude of products. Grain-based resins have also been used. For example, U.S. Patent No. 5,665,152 issued to Bassi et al., which is incorporated by reference herein, describes formulations and processing methods for grain-based protein products. Grain proteins may be prepared as resin pellets, which can then be used for many applications, including extrusion and injection molding applications. However, if the resin pellets are not used in these processes within a short time of resin production, e.g., a few weeks, the molded articles begin to show signs of rough and bumpy surfaces due to aging of the resin pellets. The strength of the injection molded articles can also decrease. These problems increase with time after resin production, which necessitates the use of the resin pellets shortly after production.

[0003] Thus, it would be a valuable contribution to the art to provide grain protein-based resin formulations having improved aging properties for use in shaped, molded, and extruded objects.

1

SUMMARY OF THE INVENTION

[0004] In one aspect, the resin aging problem is addressed by incorporating a certain amount of a shelf-stabilizing agent, such as hydrolyzed proteins, hydrolyzed protein derivatives, hydrolyzed protein/hydrolyzed protein derivative - emulsifier complexes, and mixtures thereof in the formulation for the resin pellets made by extrusion processes.

[0005] Formulation details for the making of resin pellet are also provided.

[0006] Methods of preparing the grain protein-based resin pellets and molding methods, such as injection molding, are yet further provided.

BRIEF DESCRIPTION OF THE DRAWINGS

[0007] Figure 1 illustrates a comparison of injection molded pet chew products made from aged and non-aged wheat gluten based resin pellets.

DETAILED DESCRIPTION OF THE INVENTION

[0008] The term "%" as employed throughout the specification and claims refers to weight percent unless otherwise specifically noted in the text.

[0009] Broadly speaking, one method of the invention first comprises the step of providing a formulation especially designed to have melt flow and rheological properties allowing the formulation to be processed using conventional plastics forming equipment. This formulation is then heated under moderate temperature conditions, usually with shear, to create a substantially homogeneous and flowable formulation. The most common way of making the resin pellets is by extrusion processes. Both single and twin-screw extruders can be used, with more preference being given to twin-screw extrusion due to better mixing and pumping action. The resin pellets can then be formed into desired articles using injection molding, extrusion or other forming equipment. Very importantly, the resin formulation may be prepared for later use as a substantially homogeneous and flowable product with the avoidance of any substantial heat denaturation of the grain protein. During the formation of the final desired articles, the substantially undenatured protein is denatured. Thus, in the context of injection molding, the preferred temperature conditions of molding are selected to assure essentially complete protein denaturation. It is also the case that certain water soluble denatured proteins, such as soy proteins,

may be used in the resins, in which case the resin may be further denatured or cured by the injection molding, extrusion or other forming equipment.

[0010] The resin aging problem is associated with chemical and/or conformational change in the resin pellets. It has been discovered that shelf life of the resin formulations may be significantly extended by formulating an undenatured grain-based protein with a certain amount of shelf stabilizing agent, such as hydrolyzed proteins, hydrolyzed protein derivatives, hydrolyzed protein/hydrolyzed protein derivative - emulsifier complexes, and mixtures thereof.

plasticized state undergoes heat treatment that results in the protein losing its viscoelasticity or viscous flow (melt flow) property. The melt flow viscosity increases as
the degree of heat denaturation increases. There is little or no melt flow property if
the protein is completely heat denatured. For example, corn gluten meal coming out
of the dryer of a wet milling process is severely heat denatured and, therefore, is not
useful in the pellet resin formulation to provide useful rheological properties for
purposes of the present invention. On the other hand, commercially available vital
wheat gluten is processed to provide minimum heat denaturation and, in combination
with the shelf-stabilizing agents, is a suitable grain protein source for the present
invention. Commercially available soy protein products may possess varying degrees
of heat denaturation resulting from their preparation; however, most possess good
melt flow properties with adequate plasticization due to the water solubility of soy
protein.

[0012] In more detail, the preferred grain protein-based formulation includes from about 20% to about 80% by weight grain protein, and preferably from 30% to 75%. Although a variety of grain proteins can be employed, most preferably the protein is selected from the group consisting of soy protein, wheat gluten, corn gluten and mixtures thereof. In preferred forms, the grain protein has substantially no heat denaturation and as used is naturally occurring. Normally, for reasons of economy and ease of formulation, the grain protein is provided as a part of a mixture which would typically include other optional ingredients such as starch, lipids, bran and combinations thereof. For example, soy meals, concentrates and isolates could be used, as well as various commercial grades of wheat and corn gluten. When such

mixtures are used, typically they would provide at least about 50% by weight of the desired grain protein, and more preferably at least about 75% by weight thereof.

[0013] The most important aspect of the present invention is to address the resin pellet aging problem by incorporating in the formulation a shelf-stabilizing agent, such as hydrolyzed proteins, hydrolyzed protein derivatives, hydrolyzed protein/hydrolyzed protein derivative - emulsifier complexes, or mixtures thereof.

[0014] The hydrolyzed proteins to be employed in the invention may, for example, include hydrolyzed wheat gluten, hydrolyzed soy protein, hydrolyzed corn gluten, hydrolyzed potato protein, hydrolyzed rice protein, hydrolyzed gelatin protein, hydrolyzed collagen, hydrolyzed casein, hydrolyzed whey protein, hydrolyzed milk protein, hydrolyzed egg white, hydrolyzed egg yoke, hydrolyzed whole egg, hydrolyzed chicken liver, hydrolyzed pork liver, hydrolyzed beef liver, hydrolyzed fish liver, hydrolyzed meat protein of any source, hydrolyzed fish, hydrolyzed blood plasma, and mixtures thereof. Preferred protein hydrolysates are hydrolyzed wheat gluten, hydrolyzed soy protein, hydrolyzed liver proteins. The hydrolyzed protein is generally present in the range of from about 0.5% to about 25% by weight of the formulation. Preferred for the practice of the invention is a hydrolyzed protein of from about 1.5 to about 20% by weight. Particularly preferred for the practice of the invention is a hydrolyzed protein amount of from about 2% to about 15 % by weight.

[0015] To best use the hydrolyzed proteins to address the aging problem of resin pellets, it is preferable to have the number average molecular weight and weight average molecular weight of the hydrolyzed protein component in the practice of the present invention, less than 10,000 and 20,000 Daltons, respectively.

[0016] Hydrolyzed proteins may be prepared by any means. Typically, enzymatic hydrolysis or acid hydrolysis is employed. Preferred for the practice of the present invention is enzymatic hydrolysis. The hydrolysate is typically adjusted to a pH of 4-7.5 using NaOH, KOH, Ca(OH)₂, and the like, before spray or flash drying the product.

[0017] Examples of hydrolyzed protein derivatives include reaction products of protein hydrolysates with other chemicals or low molecular weight polymer or oligomer ingredients. The reaction products contain a hydrolyzed protein moiety and a derivative portion. The amount of hydrolyzed protein in the derivative

reaction products may range from about 0.5 % to about 50 % depending on the reaction chemicals used. Examples are reaction products of hydrolyzed protein with anhydride, ethylene oxide, propylene oxide, fatty acid derivatives, reducing sugars, maltodextrin, oligosaccharides, dextrin, and the like.

[0018] The amount of hydrolyzed protein derivatives to be employed in the formulation may be from about 1% to about 25 %. Preferred for the practice of the invention is a hydrolyzed protein derivatives of from about 1.5 to about 20 %. Particularly preferred for the practice of the invention is a hydrolyzed protein derivative amount of from about 2 to about 15 %.

[0019] The hydrolyzed protein/hydrolyzed protein derivatives—emulsifier complex of the present invention may be prepared from hydrolyzed protein moieties and hydrolyzed protein derivative moieties bonded physically with emulsifiers.

Suitable emulsifiers to be used in the present invention include hydrolyzed vegetable oil, hydrolyzed animal fat, hydrolyzed lecithin and their salt forms, hydrolyzed lecithin modified further by ethylene oxide and propylene oxide, ethoxylated monoand diglycerides, diacetyl tartaric acid ester of mono-diglycerides, sugar esters of mono- and diglycerides, propylene glycol mono—and diesters of fatty acids, calcium stearoyl-2-lactylate, lactylic stearate, sodium stearoyl fumarate, succinylated monoglyceride, sodium stearoyl-2-lactylate, polysorbate 60, or any other emulsifier that contains both hydrophobic and hydrophilic portions in the structure, and mixtures thereof. The amount of emulsifiers in the complex is from about 10-30 % by weight of the complex.

[0020] The amount of hydrolyzed protein/hydrolyzed protein derivatives-emulsifier complex to be employed in the formulation for resin production may be from about 1 to about 25 %. Preferred for the practice of the invention is a hydrolyzed protein/hydrolyzed protein derivatives-emulsifier complex of from about 1.5 to about 20 %. Particularly preferred for the practice of the invention is a hydrolyzed protein/hydrolyzed protein derivatives-emulsifier complex amount of from about 2 to about 15 %.

[0021] The formulation of resin pellets may also contain from about 10-40% plasticizers in the starting formulations, and more preferably from about 10-35% by weight. The preferred class of plasticizers include those selected from the group

consisting of, glycerol, diglycerol, propylene glycol, triethylene glycol, urea, sorbitol, mannitol, maltitol, hydrogenated corn syrup, polyvinyl alcohol, polyethylene glycol, and mixtures thereof. The most preferred plasticizer is glycerol.

- [0022] The extrudable formulations of the invention may also include a minor amount of water, up to 14 % by weight, more preferably up to about 12 % by weight, and most preferably from about 2-10 % by weight. The presence of excess water leads to a sticky, stretchy extrudate unsuited for use in the formation of solid non-edible products. The moisture content in the resin pellets is preferably controlled from about 5-12 %.
- [0023] The formulation of resin pellets may also contain from about 0.5% to 5% lubricants. The presence of lubricants helps extrusion process and molding operation for ease of melt flow and melt temperature control. The lubricants may include glycerol mono/di-stearate, hydrolyzed lecithin and derivatives, fatty acid and derivatives. The preferred lubricant is glycerol monostearate.
- [0024] The formulation of resin pellets may also contain from about 0.5% to 3 % mold release agents. The presence of such releasing agent prevents the parts or articles from sticking to the molding surface or processing surface in general. The mold release agents may be magnesium stearate, calcium stearate, barium stearate, or other alkaline earth metal fatty acid agents. A particularly preferred mold release agent is magnesium stearate.
- [0025] The formulation of resin pellets may also contain from about 0.5% to 5% reducing agent. The reducing agent cleaves the disulfide bonds in the grain protein. This drastically improves the flow and mixing of the grain protein in the processing equipment, rendering the overall formulation more suitable for use therein. The reducing agent is preferably present in a minor amount of at least about 0.01% by weight, and more preferably from about 0.05-3 % by weight, where these weights are based upon the total amount of grain protein being taken as 100 % by weight. The reducing agents are advantageously selected from the group consisting of the alkali metal and ammonium sulfites, bisulfites, metabisulfites and nitrites, and mercaptoethanol, cysteine, cysteamine, sulfur dioxide, ascorbic acid and mixtures thereof. A particularly preferred reducing agent is sodium metabisulfite.

[0026] Normally, the reducing agent is simply added to the other components of the formulation prior to or as a part of the extrusion process. Alternately, the reducing agent can be used to preliminarily treat the selected grain protein(s) prior to preparation of the starting formulation. Thus, in the case of gluten products (wheat and corn gluten), the reducing agent may be initially added to obtain a modified gluten product which then is employed as a part of the extrusion formulation. In any case, the reducing agent is preferably used in an effective amount to cleave from about 5-100 % of the disulfide bonds in the grain protein.

A number of other ingredients can also be used in the starting [0027] extrusion formulations. Those optional ingredients may include: (1) fillers such as native or chemically modified starches in their granular form (wheat starch, corn starch, potato, rice, tapioca starches, and mixtures thereof, chemical modifications being oxidation, acetylation, carboxymethylation, hydroxyethylation, hydroxypropylation, and alkylation), calcium carbonate, heat denatured animal or vegetable protein granules or powder, vegetable powder, granules or special shapecuts, rice flour, wheat flour, corn gluten meal, fibers (cellulose fiber, micro-crystalline fiber, soluble fibers, wheat bran, soy bean fiber, corn grit fiber); (2) pigments (titanium dioxide, carbon black, talc, calcium carbonate); (3) coloring agents (azo dyes, chlorophyll, xanthophyll, carotene, indigo, all the synthetic colors, natural coloring agents); (4) foaming agents (sodium bicarbonate, N₂ and CO₂), and (5) other special effect ingredients such as breathe and dental cleaning ingredients. These optional ingredients may, for example, provide from about 0.001% to 75 % by weight of the resin pellets.

[0028] The formulations of the invention can be formed into pellets which can later be used in molding equipment or shaped by various methods, as illustrated in U.S. Pat. No. 5,665,152. For example, such pellets may be formed by extrusion, using either single or twin screw extruders. However, it is important to maintain the temperature of the material within the extruder barrel below about 95°C to avoid heat denaturation of the matrix protein content of the formulation. Extruded pellets of this character would generally be maintained in closed containers and would have moisture content ranging from about 5 to about 12 %.

[0029] The formulations of the present invention may be shaped into any desired object. Further, the formulations may be shaped or molded using injection molding. The melt temperature inside the barrel of the injection molder should be maintained to a level of up to about 90°C, and more preferably up to about 65°C. However, the mold itself would normally be heated to a temperature of from about 120 to about 180°C, in order to denature the grain protein fraction of the formulation introduced into the injection mold. The other parameters of injection molding such as cycle time (ranging from a few seconds to a few minutes) are as employed in the art.

[0030] Suitable products to be prepared using the grain protein-based resin formulations of the present invention include, for example, pet chew treats, edible products and biodegradable products in general.

[0031] The following examples illustrate the specific formulations and methods of preparing the resin pellets and molded articles.

EXAMPLES

SOURCES AND IDENTITY OF MATERIALS

[0032] Vital Wheat Gluten is a commercially available wheat gluten made by a flash drying process. Wheat gluten can also be made by spray drying so long as the proteins are not denatured and lose visco-elasticity or other viscous properties after hydration.

[0033] MidsolTM is a trademark of MGP Ingredients, Inc.

[0034] Solka-Floc™ is a trademark of International Fiber Corporation, and is a cellulose fiber.

[0035] PanodanTM is a trademark of Danisco, and is a stearate derivative.

[0036] OptimizorTM is a trademark of Applied Food Biotechnology, Inc., and is hydrolyzed liver protein derivatives with maltodextrin and complexed with hydrolyzed animal fat/vegetable oil.

EXAMPLE 1

[0037] Table 1 shows a resin formulation containing 8 % hydrolyzed wheat gluten protein (HWG 2009, manufactured by MGP Ingredients, Inc.) in the formulation. The resin pellets were prepared using a 85 mm twin screw extruder

(TX-85 manufactured by Wenger) with a hot face die cutter. The powder liquid mix in the extruder is mixed at a melt temperature no more than 95°C to avoid protein heat denaturing. After the pellet is cut at the die face, the resin pellets are pneumatically transferred to a cooler and packaged.

[0038] The presence of 8 % hydrolyzed wheat gluten helps increase the shelf life of the resin pellets checked after 4 months. The molded articles exhibited the same characteristics, both appearance and physical properties, as the one molded right after the resin is produced.

TABLE 1
Resin formulation with 8 % hydrolyzed wheat gluten

Ingredient	Weight Percent
Midsol ™ HWG 2009	8
Hydrolyzed wheat gluten	
vital wheat gluten	66
glycerol monostearate	2.0
magnesium stearate	0.9
glycerine	18.5
water	2.5
sodium metabisulfite	0.1
Solka-floc™ 900	2.0
cellulose fiber	

EXAMPLE 2

[0039] Table 2 shows a formulation containing 5 % hydrolyzed wheat gluten protein-emulsifier complex in the formulation. The hydrolyzed protein—emulsifier complex was prepared by adding Panodan™ SDK emulsifier (provided by Danisco) into the HWG 2009 protein dispersion and spray dried. The hydrolyzed protein—emulsifier complex contains 25 % Panodan™ SDK (emulsifier) in the spray dried product.

[0040] The resin pellets were prepared using a 85 mm twin screw extruder (TX-85 manufactured by Wenger) with a hot face die cutter. The powder liquid mix

in the extruder was mixed at a melt temperature of no more than 95 °C to avoid protein heat denaturation. After cutting at the die face, the resin pellets were pneumatically transferred to a cooler and packaged.

[0041] The presence of 5 % hydrolyzed wheat gluten-emulsifier complex helped increase the shelf life of the resin pellets checked after 5 months. The molded articles exhibited the same characteristics, both appearance and physical properties, as the one molded right after the resin is produced.

TABLE 2

Resin formulation with 5 % hydrolyzed wheat gluten-emulsifier complex

Ingredient	Weight Percent
Hydrolyzed wheat gluten – emulsifier complex Prepared as described above	5
vital wheat gluten	59
glycerol monostearate	2.0
magnesium stearate	0.9
glycerine	18.5
water	2.5
sodium metabisulfite	0.1
Solka-floc [™] 900	2.0
cellulose fiber	
Wheat flour	10

EXAMPLE 3

[0042] Table 3 shows a formulation containing 3.5 % liver digest (OptimizorTM CHX-base, manufactured by Applied Food Biotechnology, Inc) in the formulation. CHX-Base liver digest is a form of hydrolyzed protein, hydrolyzed protein derivative and emulsifier complex where the protein is a poultry liver, and the emulsifier is hydrolyzed animal fats. The hydrolyzed protein derivatives are reaction products of hydrolyzed protein with maltodextrin for aroma enhancement. The resin pellets was prepared using a 85 mm twin screw extruder (TX-85 manufactured by Wenger) with a face die cutter. The powder liquid mix in the extruder is mixed at a melt temperature no more than 95°C to avoid protein heat denaturing. After die face pellet, the resin pellets are pneumatically transferred to a cooler and packaged.

[0043] The presence of 3.5 % CHX-Base liver digest assists in increasing the shelf life of the resin pellets checked after 5 months. The molded articles exhibited the same characteristics, both appearance and physical properties, as the one molded right after the resin is produced.

1 1

TABLE 3
Resin formulation with 3.5 % liver digest

Ingredient	Weight Percent
Optimizor™ CHX-Base	3.5
vital wheat gluten	70.5
glycerol monostearate	2.0
magnesium stearate	0.9
Propylene glycol	18.5
water	2.5
sodium metabisulfite	0.1
Solka-floc™ 900	2.0
(cellulose fiber)	

EXAMPLE 4

[0044] The resin pellets made according to example 1-3 are injection molded with an injection molding machine. Typical barrel temperature settings of the injection molding machine are: 75°C (tip end), 70°C, 60°C, 60°C (feed end). The mold temperature is set at 145°C.